Structural-functional connectivity deficits of neocortical circuits in the Fmr1<sup>−/−</sup> mouse model of autism

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Fragile X syndrome (FXS), the most common inherited form of intellectual disability disorder and a frequent cause of autism spectrum disorder (ASD), is characterized by a high prevalence of sensory symptoms. Perturbations in the anatomical connectivity of neocortical circuits resulting in their functional defects have been hypothesized to contribute to the underlying etiology of these disorders. We tested this idea by probing alterations in the functional and structural connectivity of both local and long-ranging neocortical circuits in the Fmr1<sup>−/−</sup> mouse model of FXS.

To achieve this, we combined in vivo ultrahigh-field diffusion tensor magnetic resonance imaging (MRI), functional MRI, and viral tracing approaches in adult mice. Our results show an anatomical hyperconnectivity phenotype for the primary visual cortex (V1), but a disproportional low connectivity of V1 with other neocortical regions. These structural data are supported by defects in the structural integrity of the subcortical white matter in the anterior and posterior forebrain. These anatomical alterations might contribute to the observed functional decoupling across neocortical regions. We therefore identify FXS as a “connectopathy,” providing a translational model for understanding sensory processing defects and functional decoupling of neocortical areas in FXS and ASD.

RESULTS

Reduced structural integrity of the corpus callosum

To provide an initial characterization of the anatomical connectivity of the whole brain of adult Fmr1<sup>−/−</sup> mice and wild-type littermate controls,
we performed high-resolution (11.7 T) diffusion tensor imaging (DTI) permitting the identification of the corpus callosum—the major fiber bundle serving the neocortex. We calculated the fractional anisotropy (FA) values, permitting an exploration of the orientation coherence of axons in this fiber bundle (32). We found that the FA values are reduced in several regions of the corpus callosum of Fmr1<sup>−/−</sup> mice (cf. wild-type controls), suggesting diminished structural integrity of this fiber tract (Fig. 1). Specifically, FA values were decreased in the posterior part of the brain, the splenium/forceps minor (FMI) of the corpus callosum [FA(wild type)<sub>FMI</sub> = 0.427 ± 0.005; FA(Fmr1<sup>−/−</sup>)<sub>FMI</sub> = 0.391 ± 0.009; P = 0.0069], and in the anterior part of the brain, the forceps minor (FMI) of the corpus callosum [FA(wild type)<sub>FMI</sub> = 0.386 ± 0.007; FA(Fmr1<sup>−/−</sup>)<sub>FMI</sub> = 0.329 ± 0.019; P = 3.5 × 10<sup>−3</sup>]. No changes were seen in the external capsule (EC) [FA(wild type)<sub>EC</sub> = 0.347 ± 0.011; FA(Fmr1<sup>−/−</sup>)<sub>EC</sub> = 0.321 ± 0.008; not significant (n.s.)] and the central part of the corpus callosum, Genu/Body (G/B) [FA(wild type)<sub>G/B</sub> = 0.354 ± 0.003; FA(Fmr1<sup>−/−</sup>)<sub>G/B</sub> = 0.345 ± 0.007; n.s.] (Fig. 1). No significant changes were found in the measurements of other diffusion-related parameters, such as average diffusivity and first eigenvalue, either in the corpus callosum or in the neocortical areas (fig. S1). Together, these findings point to an alteration in the orientation of fibers within the corpus callosum rather than to changes in their absolute number. For example, a relative decrease of the overall long-range connectivity compared to the local/short-range connectivity could reduce the FA value.

**Alterations in the structural connectivity of the primary visual cortex**

The aforementioned results, showing an altered structural organization of the white matter in Fmr1<sup>−/−</sup> mice, suggest that the adjacent neocortical areas—in particular the frontal cortex and the primary visual cortex (V1)—may be strongly affected by these changes. Sensory information processing defects are a common feature of both FXS and ASD [for example, Lane et al. (22) and Marco et al. (33)], and functional imaging studies suggest increased activation of the neocortex of ASD subjects in response to sensory stimuli of a range of modalities (24). In addition, alterations in visual perception and fine-scale neuroanatomical defects in the visual cortex (V1) have been reported for FXS patients (34–37). Indeed, rescue of an aberrant spine phenotype in V1 of Fmr1<sup>−/−</sup> mice has been used as an end point for pharmacological rescue approaches (38). In light of these findings and our own aforementioned results, we probed the reorganization of the anatomical connectivity of V1 by mapping and quantifying neurons projecting to this region. To permit this analysis, we injected a retrograde viral tracer into V1 and subsequently quantified local and long-ranging projections [as the fraction of labeled neurons (FLN)]. To determine the precise location of the injection site, we simultaneously co-injected an anterograde variant of the same virus (39). The cell body locations of all labeled cells were then introduced as fiducial markers into a three-dimensional (3D) brain model (40) to compute the Euclidian distance in the whole brain from the injection site (Fig. 2; A and B; P < 0.0001, Mann-Whitney test), which supports the "long-range hypoconnectivity versus local hyperconnectivity hypothesis of autism" (2). However, the concept of local connectivity is, in general, poorly defined in the literature and can be expressed (i) as a factor of distance from the injection site or (ii) with respect to a functionally defined brain area. We found that the number of local connections was increased in Fmr1<sup>−/−</sup> mice according to both definitions (Fig. 2, C to F). For example, the FLN for proximal locations is significantly greater for Fmr1<sup>−/−</sup> cf. wild-type subjects [Fig. 2C; FLN(wild type)<sub>proximal</sub> = 28.52 ± 6.17%; FLN(Fmr1<sup>−/−</sup>)<sub>proximal</sub> = 53.63 ± 3.52%; P = 0.0242 using unpaired t test]. Likewise, quantifying the FLN as a function of brain area shows a significant bias toward intrinsic inputs from within V1 [Fig. 2, D to F; FLN(wild type)<sub>intrinsic</sub> = 45.79 ± 4.29%; FLN(Fmr1<sup>−/−</sup>)<sub>intrinsic</sub> = 62.28 ± 5.96%; P = 1.57 × 10<sup>−3</sup> using multiple t tests corrected for multiple comparisons using the Holm-Sidak method with α = 0.05]. When considering only the intrinsic input, we additionally found an increased local "input clustering" in Fmr1<sup>−/−</sup> mice. Specifically, more incoming projections arrive from cells belonging to the nucleus basalis of Meynert.
in proximity to the target zone (Fig. 2E; P < 0.0001, Mann-Whitney test) with a peak of input density at ~220 to 260 µm (Fig. 2F).

Functional decoupling of neocortical regions
Increased clustering has previously been shown to alter the dynamics of network activity (41) and may also serve to amplify the intrinsic signal at the expense of more distant inputs. An overrepresentation of the local information and a reduction of input from other regions would cause the target region to be more isolated and functionally decoupled from other brain areas. To test this possibility, we measured spontaneous activity using functional magnetic resonance imaging (fMRI) in head-restrained Fmr1−/− mice under light isoflurane anesthesia. The local connectivity changes impinging on the sensory circuits in Fmr1−/− mice (Fig. 3 and fig. S4) was more strongly affected than the homotypic interhemispheric connectivity (Fig. 3, B and D). However, we noted a general trend for reduced homotypic interhemispheric connectivity in the neocortical areas, with significant differences noted for the motor cortex [fig. S4; P = 0.0034 using multivariate analysis of variance (ANOVA), corrected for multiple comparisons]. Together, we provide evidence for global changes in functional connectivity in the forebrain of adult Fmr1−/− mice.

DISCUSSION
Here, we present converging evidence pointing to both large-scale and local connectivity changes impinging on the sensory circuits in Fmr1−/− mice. In particular, we found a localized reduction in FA values, which points to white matter defects suggestive of alterations in the organization of the corpus callosum. We demonstrate, at the anatomical level, a local hyperconnectivity in V1 versus a long-range hypoconnectivity affecting global inputs into V1. These structural findings are corroborated by evidence of widespread cross-area functional hypoconnectivity, affecting the sensory circuits (among others) of Fmr1−/− mice. Together, these findings lend support to the theory of "long-range hypoconnectivity versus local hyperconnectivity" in ASDs.

The local hyperconnectivity phenotype reported here for the visual cortex is consistent with recent findings from resting-state fMRI studies of adolescents with ASD (42). Although the methods used in the current
The development of quantitative biomarkers either for the evaluation of treatment response or for patient stratification is seen as a critical step toward the development of improved clinical trials for both ASD and FXS (48–50). Such markers could be used to identify the effect of a clinical treatment or provide an advanced means of selecting patients who are likely to respond to a particular treatment especially in a multifaceted disorder like ASD (22, 47, 51, 52). Neuroimaging approaches are suggested to be particularly promising because they are objectively measurable and thought to reflect the underlying neurobiology of the disorder (53). The convergence of ASD susceptibility genes and FMRP (fragile X mental retardation protein) targets on pathways and processes governing connectivity (9, 10) further suggest that these molecules might also be targeted for pharmacological rescue. The next step in the validation of our findings would be to test whether candidate drugs can influence the aforementioned quantifiable parameters of structural or functional connectivity.

Although short-range connectivity changes have been described in Fmr1<sup>−/−</sup> mice during early postnatal development (that is, 2 to 4 weeks postnatal), these alterations are generally thought to be transient in nature (14, 16, 19, 45, 54–56). Such changes have led to a theory of a critical window for cellular and structural plasticity, suggesting that intervention during this developmentally sensitive time window is crucial for this neurodevelopmental disorder (56, 57). Here, we show a connectivity phenotype present in adult Fmr1<sup>−/−</sup> mice that challenges the notion that alterations in connectivity are restricted to this early developmental window. Largely in agreement with these findings are recent studies in humans, which identified large-scale network deficits in adolescent and adult FXS patients (58, 59). Particularly compelling...
are the findings that functional connectivity affecting a number of neocortical networks was reduced (38). In addition, the integrity of certain white matter structures was also reduced based on DTI in adolescence/adulthood and childhood (60, 61). Although it is clearly beyond the scope of the current study, it is important to note that rewiring of cortical circuits is possible, even during adulthood in healthy animals (62). Moreover, therapeutic rescue of connectivity features has previously been demonstrated in adolescence/adulthood in murine models of both FXS and ASD (38, 63).

Although we have chosen to focus on connectivity changes affecting the sensory regions of the neocortex in adulthood, this is by no means the only region implicated in the pathophysiology of FXS or ASD. Indeed, even within the context of sensory information processing, a variety of brain structures may play a role in the refinement of the neocortical circuits required for the integration of sensory information (64). Previous neuroimaging studies in individuals with FXS have identified a range of brain structures with abnormal volume, including caudate nucleus, amygdala, hippocampus, and specific regions of the neocortex and cerebellum [reviewed by Lightbody and Reiss (53)]. More diverse changes have been more recently reported in both gray and white matter volumes in younger children with FXS (65), possibly reflecting refinements in imaging technology and developmental stage. For ASD in general, the findings are more diverse. Indeed, the overall pattern is one of heterogeneity, rather than defects in specific structures, reflecting perhaps differences in image analysis, varying criteria for subject inclusion, and the heterogeneity inherent within this population (22, 47, 66). With relevance to the current work, a meta-analysis of the existing structural MRI data suggests both an alteration of cortical thickness in the parietal lobes and reduced structural integrity of the corpus callosum (66). A recent study using MRI-based neuroanatomical characterization of diverse mouse models of ASD suggests that a mathematical clustering approach might be used to identify mouse models likely to respond to the same therapeutic rescue approach (67). In light of these findings, it would be interesting to determine whether mouse models co-clustering with the *Fmr1*−/− model exhibit similar patterns of long- and short-range connectivity.

### MATERIALS AND METHODS

#### Mice

Adult (9 to 12 weeks old) male *Fmr1*−/− mice (12) in a C57BL/6 background and their male wild-type littermates [as described by Zhang et al. (28)] were used in all experiments. Animals were bred at two sites, namely, the SPF (specific pathogen-free) animal facility of the Neurocentre Magendie (Bordeaux, France) and the animal facility of the Radboud University (Nijmegen, Netherlands). All experiments were performed according to the European Directive governing the use of experimental animals (2010/63/EU) and local institutional guidelines; all experiments received previous ethical approval from the Ethics Committee of Bordeaux (CE2A50; approval #5012024-A). Genotypes were determined by a polymerase chain reaction analysis of DNA extracted from tail samples. All experiments and analysis were performed with the experimenter being blind to the genotype.

#### Virus production

SAD ΔG-eGFP (RG), a retrograde variant of glycoprotein-deleted recombinant rabies virus, was produced as previously described by Haberl et al. (39). VSV G<sup>RtmC</sup>-pseudotyped SAD ΔG-mCherry, an anterograde variant of the same virus, was produced as previously described by Haberl et al. using BSR T7/5 cells (39).

#### Stereotactic injections

Stereotactic injections were performed in *Fmr1*−/− and wild-type mice at 10 to 12 weeks of age. The stereotactic injections of viral vectors were performed in isoflurane-anesthetized and head-fixed mice using a 10-μl glass syringe fitted with a 34-gauge needle or a pulled glass pipette. Injection volume and speed were controlled using a WPI Ultra Micro Pump. Viral injections were performed using a 9:1 mixture of retrograde RABV ΔG (expressing enhanced green fluorescent protein) and anterograde RABV ΔG (expressing mCherry) as described by Haberl et al. (68). mCherry expression at the site of injection was used to verify injection coordinates and determine the precise location of the injection. Injection coordinates for upper layer 5 of V1 were at anterior/posterior (A/P) 2.8 mm, lateral (L) 2.25 mm, and dorsal/ventral (D/V) 0.5 mm. A/P and L coordinates are given with respect to the bregma, whereas D/V coordinates are given with respect to the brain surface. Precise coordinates were later validated for each injection by calculating the center of mass of all anterogradely [SAD ΔG-mCherry (VSV G<sup>RtmC</sup>)]-labeled neurons in the respective brain (COM<sub>r</sub>). To verify that all injections are comparable (mapping the same area), we calculated the SD from the injection coordinates using the COM<sub>r</sub> of all analyzed injections: A/P, 2.8 ± 0.175 mm; L, 2.25 ± 0.126 mm; and D/V, 0.5 ± 0.06 mm. All distance calculations were performed for every cell to the COM<sub>r</sub> of the respective brain.

#### Mouse brain slice preparation

Mice were perfused and brains were sectioned as previously described by Haberl et al. (39). Briefly, mice were administered a lethal dose of sodium pentobarbital and then transcardially perfused with 30 ml of normal Ringer’s solution followed by 100 ml of a 4% paraformaldehyde (PFA) solution in 1× phosphate-buffered saline. The mouse brains were dissected and postfixed in 4% PFA solution and slices were cut using a vibratome (Leica). For whole forebrain sectioning, brains were immersed in 10% gelatin, postfixed 2 hours in 4% PFA, sectioned in 50-μm slices, and mounted using Prolong Gold Antifade Reagent.

#### Fluorescence microscopy and analysis

Images of the entire forebrain were acquired using a scanning mosaic wide-field fluorescence acquisition system (NanoZoomer, Hamamatsu) equipped with a 20× 0.75 numerical aperture objective. Images were acquired by scanning each section at multiple (about five to six) z-positions with an 8-μm step size. Analysis of retrogradely and locally labeled cells was performed blind to the genotype. Fiducial markers were placed manually on the cell position, and images were segmented using the mouse brain atlas (*The Mouse Brain in Stereotaxic Coordinates*) by K. B. J. Franklin and G. Paxinos, (1997). Fiducial points transformed in a 3D average brain atlas in Vaa3D software (69) were used to compute the Euclidian distance of each marker in the whole brain in *xyz* directions from the injection site. The brain model was previously generated from anatomical MRI scans of adult C57BL/6 mice at 16-μm spatial resolution (40). The precise coordinates of each injection site were confirmed by calculating the center of mass of cells infected by the local tracer SAD ΔG-mCherry (VSV-G). We counted a total of 7209 retrogradely labeled neurons and calculated an average distance of 973 μm to its respective COM<sub>r</sub>, (1183 μm in the wild-type mice and...
810 μm in the Fmr1<sup>−/y</sup> mice) equaling a total minimum wiring length of 6.94 m (1.22 ± 0.09 m in wild type versus 1.09 ± 0.07 m in Fmr1<sup>−/y</sup>).

**Definition.** To test the hypothesis of short-ranging hyperconnectivity, we determined the intrinsic and the proximal FLN. Retrogradely labeled neurons within V1 are intrinsic. Proximity was set as distance below 500 μm.

**Magnetic resonance imaging**

MRI measurements were performed with an 11.7-T BioSpec Avance III small animal MR system (Bruker BioSpin) equipped with an actively shielded gradient set of 600 mT/m and operated by Paravision 5.1 software. We used a circular polarized volume resonator for signal transmission and an actively decoupled mouse brain quadrature surface coil for signal reception (Bruker BioSpin). The levels of anesthesia and mouse physiological parameters were monitored following an established protocol to obtain a reliable measurement of functional connectivity (70). Briefly, during the MR experiments, low-dose isoflurane was used (3.5% for induction and ~1.5% for maintenance), slightly adjusted throughout the experiment to maintain a fast and stable breathing frequency (>130 beats/min). The mice were placed in a stereotactic device to immobilize the head. Body temperature was measured with a rectal thermometer and maintained at 37°C by a heated airflow device.

After standard adjustments and shimming, fMRI data sets were acquired using a single-shot spin-echo sequence combined with echo-planar imaging (SE-EPI) sequence. Six hundred repetitions with a repetition time (TR) of 1.8 s and an echo time (TE) of 16.9 ms were recorded for a total acquisition time of 18 min. Other imaging parameters were as follows: field of view, 25 × 25 mm; image matrix, 96 × 96; spatial resolution, 260 × 260 × 500 μm; number of slices, 9.

Diffusion of water was imaged as previously described by Zerbi et al. (32) and Harsan et al. (71). In short, 22 axial slices covering the whole brain were acquired with a four-shot SE-EPI protocol. B<sub>0</sub> shift compensation, navigator echoes, and an automatic correction algorithm to limit the occurrence of ghosts and artifacts were implemented. Encoding b factors of 0 s/mm<sup>2</sup> (b<sub>0</sub> images; 5×) and 1000 s/mm<sup>2</sup> were used and diffusion-sensitizing gradients were applied along 30 noncollinear directions in 3D space. Other imaging parameters were as follows: TR, 7.55 s; TE, 20 ms; field of view, 20 × 20 mm; image matrix, 128 × 128; spatial resolution, 156 × 156 × 500 μm; total acquisition time, 18 min.

**Functional connectivity measurements**

The fMRI data sets were processed as previously described by Zerbi et al. (70). Briefly, the data were first realigned using a least-squares method and rigid-body transformation with Statistical Parametric Mapping (SPM) mouse toolbox [SPM5, University College London (72)]. The mean SE-EPI images of each mouse were then used to generate and normalize the data into a study-specific template through linear affine and nonlinear diffeomorphic transformation [ANTS (Advanced Normalization Tools) v1.9, http://picsl.upenn.edu/ANTS/]. On the template, 17 areas were selected in the left and right hemispheres and back-transformed in each subject space using the inverse of the affine and diffeomorphic transformations. Brain regions were segmented based on an MRI atlas (40) and include the following: dorsal hippocampus, ventral hippocampus, auditory cortex, primary motor cortex, somatosensory cortex, primary visual cortex, retrosplenial cortex, piriform cortex, amygdala, pretectal area, caudate putamen, lateral geniculate nucleus, globus pallidus, parafascicular nucleus, ventral posterolateral nucleus, and ventral postero medial nucleus. In-plane spatial smoothing (0.4 × 0.4 mm) and temporal high-pass filtering (cutoff at 0.01 Hz) were applied to compensate for small across-mouse misregistration and temporal low-frequency noise using the FEAT tool of FSL [FSL 5.0 (73)]. Functional connectivity between regions of interest (ROIs) was calculated from the blood oxygen level–dependent time series after movement regression using total correlation analyses implemented in FSLNets (FSLNets, V0.3, www.fmrib.ox.ac.uk/fsi). Pearson correlation values were Fisher-transformed to Z scores for group comparisons and statistical analysis.

**Diffusion tensor MRI parameter estimation**

The calculation of the two commonly used DT-MRI parameters, mean diffusivity (MD) and FA, was performed following a protocol as previously described by Zerbi et al. (32). Briefly, the diffusion images were first realigned with SPM mouse toolbox to compensate for small movement artifacts; thereafter, the data sets were spatially normalized to a study-specific template through linear affine and nonlinear diffeomorphic transformation using ANTs. Following these preprocessing steps, the diffusion tensor was estimated for every voxel using the PATCH algorithm (74). ROIs in several white matter and gray matter areas were drawn on the template image based on an anatomical atlas (The Mouse Brain in Stereotaxic Coordinates), and the resulting FA and MD values were measured for further statistical analyses.

**Statistical analysis**

Prism 6.0e was used for analysis with statistical tests as described. All values were presented as means ± SEM unless stated otherwise. Two-tailed unpaired t tests were performed to evaluate the difference between two groups of data. Mann-Whitney test was used to compare two groups in the absence of Gaussian distribution. Two-sample Kolmogorov-Smirnov test was used to test for the equality of distributions. Multiple t tests corrected for multiple comparison using the Holm-Sidak method and multivariate ANOVA corrected for multiple comparison using Bonferroni’s method were used for the analysis of more than two parameters. The specific details are provided in the text and in each figure legend.

**SUPPLEMENTARY MATERIALS**

Supplementary material for this article is available at http://advances.sciencemag.org/cgi/content/full/1/e1500775/DC1

Fig. S1. High-field 11.7-T DT-MRI measurements of the white and gray matter of adult Fmr1<sup>−/y</sup> and wild-type littermate mice.

Fig. S2. Tracing of the input to V1.

Fig. S3. Schematic representation of the experimental strategy for the 3D anatomical registration of projection neurons into V1.

Fig. S4. Functional connectivity matrix of wild-type and Fmr1<sup>−/y</sup> mice.

Movie S1. Slice view and 3D view, illustrating the 3D mouse brain model with the combined positions of all retrogradely (input; green) and anterogradely (local; red) labeled neurons in wild-type mice.

Movie S2. Slice view and 3D view, illustrating the 3D mouse brain model with the combined positions of all retrogradely (input; green) and anterogradely (local; red) labeled neurons in Fmr1<sup>−/y</sup> mice.

REFERENCES AND NOTES


Fragile X mice develop sensory hyperreactivity to auditory stimuli.


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